In the Claims

 (Previously presented) A coating for an implantable medical device, the coating comprising a first region having a drug incorporated therein, and a second region disposed over the first region,

wherein the second region comprises a polymer and a substance having the melting temperature within the range between about 32 °C and 40°C for modifying the rate of release of the drug, the polymer having in a dry state a glass transition temperature within a range of between about 35°C and about 50°C.

wherein the polymer in the dry state contains less than about 1 mass % of water, and wherein when the body temperature of a patient in which the device is implanted rises to a temperature above the patient's normal body temperature, the morphology of coating changes so as to change the release rate of the drug in the coating.

- (Original) The coating of Claim 1, wherein the implantable medical device is a stent.
 - 3. (Original) The coating of Claim 1, wherein the drug is an anti-inflammatory drug.
- (Original) The coating of Claim 1, wherein the polymer comprises acrylic polymers, non-acrylic polymers, or blends thereof.
- 5. (Withdrawn) The coating of Claim 4, wherein the acrylic polymers are selected from a group consisting of poly(tert-butyl acrylate), poly[3-chloro-2,2-bis(chloromethyl) propyl acrylate], poly(cyanobenzyl acrylate), poly(2-methoxycarbonylphenyl acrylate), poly(3-methoxycarbonylphenyl acrylate), poly(4-ethoxycarbonylphenyl acrylate), poly(hexadecyl acrylate), poly(3-dimethylaminophenyl acrylate), poly(p-tolyl acrylate), poly(n-butyl acrylamide), poly(iso-decyl acrylamide), poly(cafluoropentyl methacrylate), poly(3,3-

dimethylbutyl methacrylate), isotactic poly(methyl methacrylate), poly(n-propyl methacrylate), isotactic poly(ethyl chloroacrylate), poly(ethyl fluoromethacrylate), and blends thereof.

- 6. (Original) The coating of Claim 4, wherein the non-acrylic polymers are selected from a group consisting of, poly(2-cyclohexylethylethylene), atactic poly(iso-propylethylene), poly(1,1,2-trimethylethylene), poly(4,4 dimethylpentylethylene), poly(2,2,2-trifluoroethoxytrifluoroethylene), poly(4-methoxybenzoylethylene), poly(3,4-dimethoxybenzoylethylene), poly(vinyl fluoride), poly(cyclopentanoyloxyethylene), 60% syndiotactic poly(formyloxyethylene), poly[4-(sec-butoxymethyl) styrene], poly(4-butoxystyrene), and blends thereof.
 - 7. (Withdrawn) The coating of Claim 4, wherein the acrylic polymers have formula

$$\begin{array}{c|cccc} X & X' & X'' \\ & & | & | & | \\ -[CH_2-C]_{m^-}[CH_2-C]_{n^-}[CH_2-C]_{p^-} \\ & | & | & | & | \\ CO-Z & CO-Z' & CO-Z'' \end{array}$$

wherein:

X, X', and X" is each, independently, a hydrogen atom or an alkyl group, such as methyl group;

Z, Z', and Z" is each, independently, a substituted or unsubstituted amino group or an alkoxy group OR,OR', and OR", where R, R' and R" is each, independently, a C₁ to C₁₂ straight chained or branched aliphatic radical; and

"m," "n," and "p" is each an integer, where m>0, $n\geq 0,$ and $p\geq 0.$

 (Previously presented) The coating of Claim 1, wherein the polymer has the melting temperature above about 50 °C. (Previously presented) A coating for an implantable medical device, comprising a
polymer, a drug incorporated therein, and a substance having the melting temperature within the
range between about 32 °C and 40°C,

wherein when the body temperature of a patient in which the device is implanted rises to a temperature above the patient's normal body temperature, the morphology of the coating changes so as to change the release rate of the drug in the coating.

- (Original) The coating of Claim 9, wherein the implantable medical device is a stent
- 11. (Previously presented) The coating of Claim 9, wherein the polymer has a glass transition temperature of the polymer in a dry state is about 37°C, wherein the polymer in the dry state contains less than about 1 mass % of water.
- (Original) The coating of Claim 9, wherein the polymer comprises acrylic polymers, non-acrylic polymers, or blends thereof.
- 13. (Withdrawn) The coating of Claim 12, wherein the acrylic polymers are selected from a group consisting of poly(tert-butyl acrylate), poly[3-chloro-2,2-bis(chloromethyl) propyl acrylate], poly(cyanobenzyl acrylate), poly(2-methoxycarbonylphenyl acrylate), poly(3-methoxycarbonylphenyl acrylate), poly(4-ethoxycarbonylphenyl acrylate), poly(hexadecyl acrylate), poly(3-dimethylaminophenyl acrylate), poly(p-tolyl acrylate), poly(n-butyl acrylamide), poly(iso-decyl acrylamide), poly(octafluoropentyl methacrylate), poly(3,3-dimethylbutyl methacrylate), isotactic poly(methyl methacrylate), poly(n-propyl methacrylate), isotactic poly(ethyl chloroacrylate), poly(ethyl fluoromethacrylate), and blends thereof.
- (Withdrawn) The coating of Claim 12, wherein the acrylic polymers have a formula

$$\begin{array}{c|cccc} X & X' & X'' \\ & & & | & & | \\ -[CH_2-C]_{m^-}[CH_2-C]_{n^-}[CH_2-C]_{p^-} \\ & & & | & & | \\ CO-Z & CO-Z' & CO-Z'' \end{array}$$

wherein:

X, X', and X'' is each, independently, a hydrogen atom or an alkyl group, such as methyl group;

Z, Z', and Z" is each, independently, a substituted or unsubstituted amino group or an alkoxy group OR,OR', and OR", where R, R' and R" is each, independently, a C₁ to C₁₂ straight chained or branched aliphatic radical; and

"m," "n," and "p" is each an integer, where m > 0, $n \ge 0$, and $p \ge 0$.

- 15. (Original) The coating of Claim 12, wherein the non-acrylic polymers are selected from a group consisting of, poly(2-cyclohexylethylethylene), atactic poly(*iso*-propylethylene), poly(1,1,2-trimethylethylene), poly(4,4 dimethylpentylethylene), poly(2,2,2-trifluoroethoxytrifluoroethylene), poly(4-methoxybenzoylethylene), poly(3,4-dimethoxybenzoylethylene), poly(vinyl fluoride), poly(cyclopentanoyloxyethylene), 60% syndiotactic poly(formyloxyethylene), poly[4-(*sec*-butoxymethyl) styrene], poly(4-butoxystyrene), and blends thereof.
 - 16. (Original) The coating of Claim 9, wherein the drug is an anti-inflammatory drug.
- 17. (Withdrawn) A method of coating an implantable medical device, comprising depositing a first layer on the device, the first layer including a drug incorporated therein, and depositing a second layer over the first layer, the second layer comprising a polymer for modifying the rate of release of the drug, wherein the polymer has a glass transition temperature

in a dry state within a range of between about 35°C and about 50°C, wherein the polymer in the dry state contains less than about 1 mass % of water.

- (Withdrawn) The method of Claim 17, wherein the implantable medical device is a stent.
- (Withdrawn) The method of Claim 17, wherein the therapeutic agent is an antiinflammatory drug.
- (Withdrawn) The method of Claim 17, wherein the polymer comprises acrylic polymers, non-acrylic polymers, or blends thereof.
- 21. (Withdrawn) The method of Claim 20, wherein the acrylic polymers are selected from a group consisting of poly(tert-butyl acrylate), poly[3-chloro-2,2-bis(chloromethyl) propyl acrylate], poly(cyanobenzyl acrylate), poly(2-methoxycarbonylphenyl acrylate), poly(3-methoxycarbonylphenyl acrylate), poly(4-ethoxycarbonylphenyl acrylate), poly(hexadecyl acrylate), poly(3-dimethylaminophenyl acrylate), poly(p-tolyl acrylate), poly(n-butyl acrylamide), poly(iso-decyl acrylamide), poly(octafluoropentyl methacrylate), poly(3,3-dimethylbutyl methacrylate), isotactic poly(methyl methacrylate), poly(n-propyl methacrylate), isotactic poly(ethyl chloroacrylate), poly(ethyl fluoromethacrylate), and blends thereof.

22. (Withdrawn) The method of Claim 20, wherein the acrylic polymers have formula

$$\begin{array}{c|cccc} X & X' & X'' \\ & & & | & & | \\ -[CH_2-C]_m-[CH_2-C]_n-[CH_2-C]_p- \\ & & & | & & | \\ CO-Z & CO-Z' & CO-Z' \\ \end{array}$$

wherein:

X, X', and X" is each, independently, a hydrogen atom or an alkyl group, such as methyl group;

Z, Z', and Z" is each, independently, a substituted or unsubstituted amino group or an alkoxy group OR,OR', and OR", where R, R' and R" is each, independently, a C₁ to C₁₂ straight chained or branched aliphatic radical; and

"m," "n," and "p" is each an integer, where m > 0, $n \ge 0$, and $p \ge 0$.

- 23. (Withdrawn) The method of Claim 20, wherein the non-acrylic polymers are selected from a group consisting of, poly(2-cyclohexylethylethylene), atactic poly(*iso*-propylethylene), poly(1,1,2-trimethylethylene), poly(4,4 dimethylpentylethylene), poly(2,2,2-trifluoroethoxytrifluoroethylene), poly(4-methoxybenzoylethylene), poly(3,4-dimethoxybenzoylethylene), poly(vinyl fluoride), poly(cyclopentanoyloxyethylene), 60% syndiotactic poly(formyloxyethylene), poly[4-(*sec*-butoxymethyl) styrene], poly(4-butoxystyrene), and blends thereof.
- 24. (Withdrawn) The method of Claim 17, wherein the polymer has the melting temperature above about 50°C, and additionally including a substance having the melting temperature within the range between about 32 °C and 40°C.